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- (54) Title: **FUEL TANKS AND FUEL TRANSPORT LINES**

(57) Abstract: A fuel tank assembly comprises a fuel tank having a wall with an outer surface and an inner surface, an elongated single or multi-walled tubular body having a first open end and a second open end, the first open end extending outwardly through an opening in the tank wall, and the second open end extending inwardly into the tank and bonded to the tank wall along the periphery of the tank wall opening by an adhesive which bonds to low surface energy plastic materials, the adhesive providing a fuel vapor-tight seal at the interface between the tubular body and tank wall opening, the fuel tank and the tubular body having fuel barrier property.

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FUEL TANKS AND FUEL TRANSPORT LINES

The present invention relates to plastic fuel tanks and fuel transport lines.

Currently, the attachments of fuel transport lines to fuel tanks are an area of concern due to cost of the assembly and the excessive emissions of fuel vapor into the atmosphere.

Plastic fuel tanks for automobiles are commonly produced by blow molding process, such as extrusion blow molding, that is, by extruding a parison into an open mold, closing the mold and blow molding the parison. Extrusion blow molding is a well known process. See, for example, H. G. Fritz "Extrusion Blow Molding," *Plastics Extrusion Technology*, Edited by Friedhelm Hensen, Hanser Publishers, pp.363-427.

Plastic fuel tanks can also be made by forming or casting a single unit or can be made by joining two or more sections into a finished unit. Further, the tanks can be formed having ports for sensor installation and for fuel inlet tubes.

Plastic fuel tanks currently use injection molded high density polyethylene (HDPE) nipples that are hot plate welded or spun-welded to the outer tank wall. A single or multi-walled tube is forced-fit over the nipple and fastened with a mechanical fastener, that is steel strap. The outer end of the tube has, or may have, a quick release fitting that attaches to the steel or polymer tubing that transports the fuel to the engine compartment. The injection molded nipple does not have fuel barrier property and, therefore, fuel vapors can permeate through the nipples. In addition, fuel vapor can

be lost through the interface between the single or multi-walled tube, the nipple, and the outer wall of the tank.

It would be desirable to provide an assembly of
35 a fuel tank and fuel transport lines which do not release
fuel vapors to the environment.

In a first aspect, the present invention is a
fuel tank assembly comprising a fuel tank having a wall
with an outer surface and an inner surface, an elongated
40 single or multi-walled tubular body having a first open
end and a second open end, the first open end extending
outwardly through an opening in the tank wall, and the
second open end extending inwardly into the tank and
bonded to the tank wall along the periphery of the tank
45 wall opening by an adhesive with some barrier properties,
the adhesive providing a fuel vapor-tight seal at the
interface between the tubular body and tank wall opening,
the fuel tank and the tubular body comprising a polymer
having fuel barrier property.

50 In a second aspect, the present invention is a
fuel tank assembly comprising a fuel tank having a wall
with an outer surface and an inner surface, a fuel
transport line having a first open end and a second open
end, the first open end extending outwardly through an
55 opening in the tank wall, and the second open end
extending inwardly into the tank and bonded to the tank
wall along the periphery of the tank wall opening by an
adhesive with some barrier properties, the adhesive
providing a fuel vapor-tight seal at the interface between
60 the fuel transport line and tank wall opening, the fuel
tank and the fuel transport line comprising a polymer
having fuel barrier property.

Fig. 1 is a cross sectional view of a single or multi-walled tubular body which is adapted to be bonded to
65 a conventional plastic fuel tank.

Fig. 2 is a cross sectional view of an alternative embodiment of the single or multi-walled tubular body shown in Fig. 1.

Fig. 3 is a cross sectional view of an assembly
70 comprising the single or multi-walled tubular body shown in Fig. 1 bonded to the inside surface of a fuel tank.

Fig. 4 is a cross sectional view of an assembly comprising a fuel transport line bonded to the inside surface of a fuel tank.

75 Referring to Figs. 1, 2 and 3, there is shown a single or multi-walled tubular body 10 with a first end 11 and a second end 12. Spaced from second end 12 is a radially and outwardly extending fluted surface 13 and a raised surface 15. Extending from one side of fluted
80 surface 13 to the nearest side of raised surface 15 is bondline 14.

In operation, tubular body 10 is pushed into (Fig. 1) or pulled through the fuel tank (Fig. 2) through a hole cut out of the wall. Fluted surface 13 and raised
85 surface 15 snap fit over tank wall 16. As shown, tubular body 10 is attached to tank wall 14 through fluted surface 13 which is bonded to tank wall 16 along bondline 14 by means of adhesive 17. Raised surface 15 holds tubular body 10 until adhesive 17 is cured to an acceptable green
90 strength. Either bondline 14 or fluted surface 13 is coated with adhesive 17. Adhesive 17 provides a fuel vapor-tight bond between tubular body 10 and tank wall 16. A quick connect may be added to the end of the tubular

body outside the tank. Quick connects are well known in
95 the art, and are described, for example in U.S. Patent
5,310,226.

Referring now to Fig. 4, the "tank end" of fuel
transport lines such as vent lines, fuel line and return
line, is provided with a radially and outwardly extending
.00 fluted surface 23 and raised surface 25. Extending from
one side of fluted surface 23 to the nearest side of
raised surface 25 is bondline 24.

As used herein, the term "tank end" refers to
the end of the vent lines, fuel line and return line which
105 is attached to the fuel tank.

To attach the fuel transport lines to a fuel
tank, each of the tank end of these lines is press-fit or
pulled through into drilled or pre-drilled holes in the
tank until the fluted surface 23 and raised surface 25
110 snap fit over tank wall 26. Fluted surface 23 is bonded
to tank wall 26 along bondline 24 by means of adhesive 27.
The fuel tank is then moved to the next assembly cell or
to a curing area.

In general, the fuel transport lines are
115 produced by extrusion, or injection molding, which is
known in the art. See, for example, U.S. Patents
6,190,154 and 6,204,312. The fluted end may be added via
compression molding or through mandrel forming operations
during a secondary operation.

120 Preferably, the plastic fuel tank, the multi-
walled tubular body and the fuel transport lines comprise
a multilayer laminate structure having one or more layers
of a low energy surface material and one or more layers of
a polymer having fuel barrier property.

125 More preferably, the plastic fuel tank, the multi-walled tubular body and the fuel transport lines comprise a three-layer laminate structure having two outer layers of a low energy surface material and a core layer of a polymer having fuel barrier property.

130 The low energy surface materials which can be employed in the practice of the present invention include polyolefins such as polyethylene and polypropylene and polytetrafluoroethylene (PTFE).

135 Polyolefins which can be employed in the practice of the present invention for preparing the multilayer laminate structure include polypropylene, polyethylene, and copolymers and blends thereof, as well as ethylene-propylene-diene terpolymers.

Preferred polyolefins are polypropylene, linear high density polyethylene (HDPE), heterogeneously-branched linear low density polyethylene (LLDPE) such as DOWLEX™ polyethylene resin (a Trademark of The Dow Chemical Company), heterogeneously branched ultra low linear density polyethylene (ULDPE) such as ATTANE™ ULDPE (a Trademark of The Dow Chemical Company); homogeneously-branched, linear ethylene/α-olefin copolymers such as TAFMER™ (a Trademark of Mitsui Petrochemicals Company Limited) and EXACT™ (a Trademark of Exxon Chemical Company); homogeneously branched, substantially linear ethylene/α-olefin polymers such as AFFINITY™ (a Trademark of The Dow Chemical Company) and ENGAGE® (a Trademark DuPont Dow Elastomers L.L.C.) of polyolefin elastomers, which can be prepared as disclosed in U.S. Patents 5,272,236 and 5,278,272; and high pressure, free radical polymerized ethylene polymers and copolymers such as low

density polyethylene (LDPE), ethylene-acrylic acid (EAA) copolymers such as PRIMACOR™ (Trademark of The Dow Chemical Company), and ethylene-vinyl acetate (EVA) copolymers such as ESCORENE™ polymers (a Trademark of Exxon Chemical Company), and ELVAX™ (a Trademark of E.I. du Pont de Nemours & Co.). The more preferred polyolefins are the homogeneously-branched linear and substantially linear ethylene copolymers with a density (measured in accordance with ASTM D-792) of 0.85 to 0.99 g/cm³, a weight average molecular weight to number average molecular weight ratio (Mw/Mn) from 1.5 to 3.0, a measured melt index (measured in accordance with ASTM D-1238 (190/2.16)) of 0.01 to 100 g/10 min, and an I10/I2 of 6 to 20 (measured in accordance with ASTM D-1238 (190/10)).

The most preferred polyolefin is a high density polyethylene. In general, high density polyethylene (HDPE) has a density of at least 0.94 grams per cubic centimeter (g/cc) (ASTM Test Method 'D-1505'). HDPE is commonly produced using techniques similar to the preparation of linear low density polyethylenes. Such techniques are described in U.S. Patents 2,825,721; 2,993,876; 3,250,825 and 4,204,050. The preferred HDPE employed in the practice of the present invention has a density of from 0.94 to 0.99 g/cc and a melt index of from 0.01 to 35 grams per 10 minutes as determined by ASTM Test Method D-1238.

Polymers having fuel barrier property which can be employed in the practice of the present invention for preparing the plastic fuel tank and the multi-walled tubular body include polyamides, polytetrafluoroethylene (PTFE), polyamides, fluoroelastomers, polyacetal homopolymers and copolymers, sulfonated and fluorinated

190 HDPE, ethylene vinyl alcohol polymers and copolymers, hydroxy-functionalized polyethers and polyesters, and branched polyesters.

Specific examples of polyamides include nylon 6, nylon 66, nylon 610, nylon 9, nylon 11, nylon 12, nylon 6/66, nylon 66/610, and nylon 6/11.

195 The single-wall tubular body which is bonded to the tank wall comprises a plastic material, such as polyethylene (also multi wall HDPE extrusions with EVOH barrier), nylon, polyester, or fluoroelastomers, or a metal material, such as steel and aluminum.

200 The tie layer, also commonly referred to as an adhesive layer, which can be employed in the practice of the present invention for preparing the multilayer structure is made of an adhesive material, such as a modified polyethylene elastomer. Preferably, the adhesive material is a maleic anhydride grafted polyethylene or 205 polypropylene such as ADMER™ (Trademark of Mitsui Petrochemicals) adhesive resin or ethylene-vinyl acetate copolymer resins such as ELVAX™ (Trademark of DuPont).

210 The adhesives which can be employed in the practice of the present invention for attaching the fuel transport lines to the fuel tank include those adhesives which bond to low energy surface plastic materials, such as the adhesive commercially known as LEA and described in an advertisement in the SPE Plastics Engineering magazine, March 2001 page 22; and adhesives comprising an 215 amine/organoborane complex, such as those described in a series of patents issued to Skoultchi (U.S. Patent Nos. 5,106,928, 5,143,884, 5,286,821, 5,310,835 and 5,376,746). These patents disclose a two-part initiator system that is

220 reportedly useful in acrylic adhesive compositions. The
first part of the two-part system includes a stable
organoborane/amine complex and the second part includes a
destabilizer or activator such as an organic acid or an
aldehyde. The organoborane compound of the complex has
three ligands which can be selected from C₁₋₁₀ alkyl groups
225 or phenyl groups. Useful amines disclosed include
octylamine, 1,6-diaminohexane, diethylamine, dibutylamine,
diethylenetriamine, dipropylenediamine, 1,3-
propylenediamine, and 1,2-propylenediamine.

230 Other adhesives which can be employed in the
practice of the present invention for attaching plastic
components to fuel tanks include those adhesives disclosed
by Zharov et al. in a series of U.S. Patents (U.S.
5,539,070; U.S. 5,690,780; and U.S. 5,691,065). These
235 patents describe polymerizable acrylic compositions which
are particularly useful as adhesives wherein
organoborane/amine complexes are used to initiate cure.
The organoboranes used have three ligands attached to the
borane atom which are selected from C₁₋₁₀ alkyl groups and
phenyl. The amine is an alkanol amine or a diamine where
240 the first amine group can be a primary or secondary amine
and the second amine is a primary amine. It is disclosed
that these complexes are good for initiating
polymerization of an adhesive which bonds to low surface
energy substrates.

245 Pocius in a series of patents (U.S. 5,616,796;
U.S. 5,6211,43; U.S. 5,681,910; U.S. 5,686,544; U.S.
5,718,977; and U.S. 5,795,657) discloses
amine/organoborane complexes with a variety of amines such
as polyoxyalkylene polyamines and polyamines which are the

250 reaction product of diprimary amines and compound having at least two groups which react with a primary amine.

The most preferred adhesive which can be employed in the practice of the present invention for attaching the fuel transport lines to the fuel tank is a
255 class of preferred amines described in copending application U.S. Serial No. 09/466321, filed December 17, 1999. These preferred amines comprise an
amine/organoborane complex wherein the organoborane is a trialkyl borane or alkyl cycloalkyl borane and the amine
260 is selected from the group consisting of (1) amines having an amidine structural component; (2) aliphatic heterocycles having at least one nitrogen in the heterocyclic ring, wherein the heterocycles may also contain one or more nitrogen atoms, oxygen atoms, sulfur atoms, or double bonds; (3) primary amines which, in addition, have one or more hydrogen bond accepting groups wherein there are at least two carbon atoms between the primary amine and the hydrogen bond accepting group, such that due to inter- or intramolecular interactions within
265 the complex, the strength of the B-N bond is increased;
270 and (4) conjugated imines.

Preferably, the trialkyl borane or alkyl cycloalkyl borane corresponds to Formula 1:

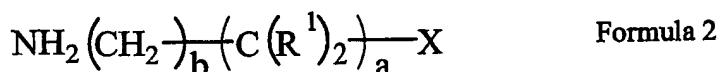


275 wherein B represents Boron; and R² is separately in each occurrence a C₁₋₁₀ alkyl, C₃₋₁₀ cycloalkyl, or two or more of R² may combine to form a cycloaliphatic ring. Preferably

R^2 is C_{1-4} alkyl, even more preferably C_{2-4} alkyl, and most preferably C_{3-4} alkyl.

280 The amine comprises a compound having a primary amine and one or more hydrogen bond accepting groups, wherein there are at least two carbon atoms, preferably at least three, between the primary amine and hydrogen bond accepting groups. Hydrogen bond accepting group means 285 herein a functional group that through either inter- or intramolecular interaction with a hydrogen of the borane-complexing amine increases the electron density of the nitrogen of the amine group complexing with the borane. Preferred hydrogen bond accepting groups include primary 290 amines, secondary amines, tertiary amines, ethers, halogen, polyethers, and polyamines.

Preferably, the amine corresponds to Formula 2:



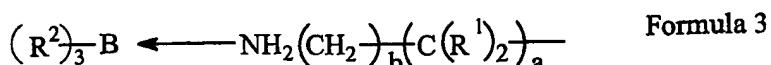
wherein:

295 R^1 is separately in each occurrence hydrogen or a C_{1-10} alkyl or C_{3-10} cycloalkyl;

300 X is hydrogen bond accepting moiety; a is an integer of 1 to 10; and b is separately in each occurrence an integer of 0 to 1, and the sum of a and b is from 2 to 10. Preferably R^1 is hydrogen or methyl. Preferably X is separately in each occurrence a hydrogen accepting moiety with the proviso that when the hydrogen accepting moiety is an amine it is a tertiary or a secondary amine. More preferably X is separately in each occurrence $-N(\text{R}^8)_e$,

305 -OR¹⁰, or a halogen wherein R⁸ is separately in each occurrence C₁₋₁₀ alkyl, C₃₋₁₀ cycloalkyl or -(C(R¹)₂)_d-W; R¹⁰ is separately in each occurrence, C₁₋₁₀ alkyl, C₃₋₁₀ cycloalkyl, or -(C(R¹)₂)_d-W; and e is 0, 1, or 2. More preferably X is -N(R⁸)₂ or -OR¹⁰. Preferably, R⁸ and R¹⁰ are
 310 C₁₋₄ alkyl or -(C(R¹)₂)_d-W, more preferably C₁₋₄ alkyl and most preferably methyl. W is separately in each occurrence hydrogen or C₁₋₁₀ alkyl or X and more preferably hydrogen or C₁₋₄ alkyl. Preferably, a is 1 or greater and more preferably 2 or greater. Preferably a is 6 or less, and most preferably 4 or less. Preferably, b is 1. Preferably, the sum of a and b is an integer 2 or greater and most preferably 3 or greater. Preferably the sum of a and b are 6 or less and more preferably 4 or less.
 315 Preferably d is separately in each occurrence an integer of 1 to 4, more preferably 2 to 4, and most preferably 2 to 3. Among preferred amines corresponding to Formula 2 are dimethylaminopropyl amine, methoxypropyl amine, dimethylaminoethylamine, dimethylaminobutylamine, methoxybutyl amine, methoxyethyl amine, ethoxypropylamine, propoxypropylamine, amine terminated polyalkylene ethers (such as trimethylolpropane tris(propylene glycol), amine-terminated)ether), aminopropylmorpholine, isophoronediamine, and aminopropylpropanediamine.

In one embodiment the preferred amine complex
 330 corresponds to Formula 3:



wherein R¹, R², X, a and b are as defined hereinbefore.

335 In another embodiment the amine is an aliphatic heterocycle having at least one nitrogen in the heterocycle. The heterocyclic compound may also contain one or more of nitrogen, oxygen, sulfur or double bonds.

340 In addition, the heterocycle may comprise multiple rings wherein at least one of the rings has a nitrogen in the ring. Preferably the aliphatic heterocyclic amine corresponds to Formula 4:



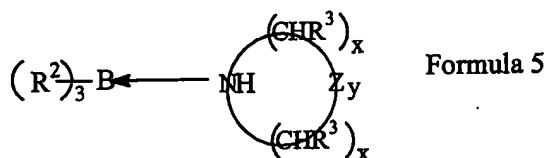
wherein:

R³ is separately in each occurrence hydrogen, a C₁₋₁₀ alkyl or C₃₋₁₀ cycloalkyl;

345 Z is separately in each occurrence oxygen or NR⁴ wherein R⁴ is hydrogen, C₁₋₁₀ alkyl, or C₆₋₁₀ aryl or alkaryl;

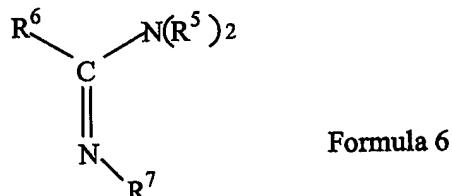
350 x is separately in each occurrence an integer of 1 to 10, with the proviso that the total of all occurrences of x should be from 2 to 10; and y is separately in each occurrence 0 or 1. Preferably, R³ is separately in each occurrence hydrogen or methyl. Preferably z is NR⁴. Preferably, R⁴ is hydrogen or C₁₋₄ alkyl, and more preferably hydrogen or methyl. Preferably 355 x is from 1 to 5 and the total of all the occurrences of x is 3 to 5. Preferred compounds corresponding to Formula 4 include morpholine, piperidine, pyrrolidine, piperazine, 1,3,3-trimethyl 6-azabicyclo[3.2.1] octane, thiazolidine, homopiperazine, aziridine, 1,4-diazabicyclo[2.2.2]octane (DABCO), 1-amino-4-methylpiperazine, and 3-pyrroline.

Complexes using aliphatic heterocyclic amines preferably correspond to Formula 5:



wherein R², R³, Z, x and y are as defined hereinbefore.

365 In yet another embodiment, the amine which is complexed with the organoborane is an amidine. Any compound with amidine structure wherein the amidine has sufficient binding energy as described hereinbefore with the organoborane, may be used. Preferable amidine
370 compounds correspond to Formula 6:

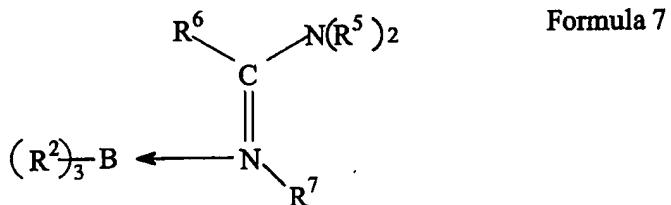


wherein:

375 R⁵, R⁶, and R⁷ are separately in each occurrence hydrogen, a C₁₋₁₀ alkyl or C₃₋₁₀ cycloalkyl; two or more of R⁵, R⁶, and R⁷ may combine in any combination to form a ring structure, which may have one or more rings. Preferably R⁵, R⁶ and R⁷ are separately in each occurrence hydrogen, C₁₋₄ alkyl or C₅₋₆ cycloalkyl. Most preferably R⁷ is H or methyl. In the embodiment where two or more of R⁵, R⁶ and R⁷ combine to form a ring structure the ring structure is
380 preferably a single or a double ring structure. Among

preferred amidines are 1,8-diazabicyclo[5.4]undec-7-ene; tetrahydropyrimidine; 2-methyl-2-imidazoline; and
385 1,1,3,3-tetramethylguanidine.

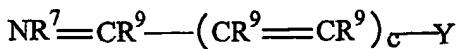
The organoborane amidine complexes preferably correspond to Formula 7:



wherein R^2 , R^5 , R^6 and R^7 are as defined earlier.

390 In yet another embodiment, the amine which is
complexed with the organoborane is a conjugated imine.
Any compound with a conjugated imine structure, wherein
the imine has sufficient binding energy as described
hereinbefore with the organoborane, may be used. The
395 conjugated imine can be a straight- or branched-chain
imine or a cyclic imine. Preferable imine compounds
correspond to Formula 8:

Formula 8



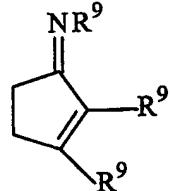
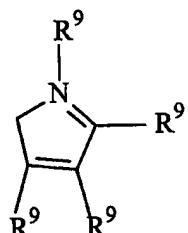
400 wherein Y is independently in each occurrence hydrogen, $N(R^4)_2$, OR^4 , $C(O)OR^4$, halogen or an alkylene group which forms a cyclic ring with an R^7 or R^9 . R^4 is hydrogen, C_{1-10} alkyl, or C_{6-10} aryl or alkaryl. Preferably R^4 is hydrogen or methyl. R^7 is as described previously. R^9 is
405 independently in each occurrence hydrogen, Y, C_{1-10} alkyl,

C_{3-10} cycloalkyl-, $(C(R^9)_2-(CR^9=CR^9)_c-Y$ or two or more of R^9 can combine to form a ring structure provided the ring structure is conjugated with respect to the double bond of the imine nitrogen; and c is an integer of from 1 to 10.

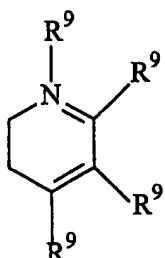
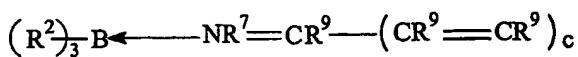
410 Preferably, R^9 is hydrogen or methyl.

Y is preferably $N(R^4)_2$, or OR^4 , or an alkylene group which forms a cyclic ring with R^7 or R^9 . Y is more preferably $N(R^4)_2$ or an alkylene group which forms a cyclic ring with R^7 or R^9 . Preferably, c is an integer of from 1 to 5, and most preferably 1. Among preferred conjugated imines useful in this invention are 4-dimethylaminopyridine; 2,3-bis(dimethylamino)-cyclopropeneimine; 3-(dimethylamine)acroleinimine; 3-(dimethylamino)methacroleinimine.

420 Among preferred cyclic imines are those corresponding to the following structures



The complexes with the conjugated imines
425 preferably correspond to Formula 9:



Formula 9

wherein R^2 , R^7, R^9 , c and Y are as defined hereinbefore.

- The molar ratio of amine compound to borane compound in the complex is relatively important. In some complexes if the molar ratio of amine compound to organoborane compound is too low, the complex is pyrophoric. Preferably the molar ratio of amine compound to organoborane compound is from 1.0:1.0 to 3.0:1.0.
- Below the ratio of 1.0:1.0 there may be problems with polymerization, stability of the complex and for adhesive uses, adhesion. Greater than a 3.0:1.0 ratio may be used although there is no benefit from using a ratio greater than 3.0:1.0. If too much amine is present, this may negatively impact the stability of the adhesive or polymer compositions. Preferably the molar ratio of amine compound to organoborane compound is from 2.0:1.0 to 1.0:1.0.

- Polymerizable compounds which may be used in the polymerization compositions of the adhesive include acrylate and/or methacrylate based compounds, with methylmethacrylate, butylmethacrylate, 2-ethylhexylmethacrylate, isobornylmethacrylate, tetrahydrofurfuryl methacrylate, and cyclohexylmethylmethacrylate as the most preferred.

Each of the polymers forming the layers of the multilayer laminate structure of the present invention may contain various additives in an amount that does not adversely affect the desired properties of the polymers.

- 455 Examples of such additives include antioxidants, ultraviolet light absorbers, thermal processing stabilizers, colorants, lubricants, flame retardants, impact modifiers, plasticizers, antistatic agents, pigments, and nucleating agents and fillers, such as
460 zeolite, talc, and calcium carbonate. The method of incorporating the additives is not critical. The additives can conveniently be added to the polymer prior to preparing the multilayer laminate structure. If the polymer is prepared in solid form, the additives can be
465 added to the melt prior to preparing the multilayer laminate structure.

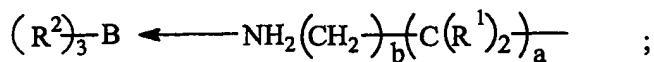
WHAT IS CLAIMED IS:

470 1. A fuel tank assembly comprising a fuel tank having a wall with an outer surface and an inner surface, an elongated single or multi-walled tubular body having a first open end and a second open end, the first open end extending outwardly through an opening in the tank wall, 475 and the second open end extending inwardly into the tank and bonded to the tank wall along the periphery of the tank wall opening by an adhesive which bonds to low surface energy plastic materials, the adhesive providing a fuel vapor-tight seal at the interface between the tubular 480 body and tank wall opening, the fuel tank and the tubular body comprising a polymer having fuel barrier property.

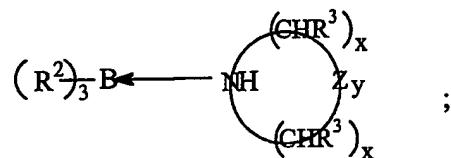
2. The fuel tank assembly of Claim 1 wherein the adhesive is an amine/organoborane complex.

3. The fuel tank assembly of Claim 2 wherein the 485 organoborane compound of the complex is a trialkyl borane or alkyl cycloalkyl borane and the amine compound is selected from the group consisting of (1) amines having an amidine structural component; (2) aliphatic heterocycles having at least one nitrogen in the heterocyclic ring, 490 wherein the heterocyclic compound may also contain one or more nitrogen atoms, oxygen atoms, sulfur atoms, or double bonds in the heterocycle; (3) primary amines which, in addition, have one or more hydrogen bond accepting groups wherein there are at least two carbon atoms between the 495 primary amine and the hydrogen bond accepting group, such that due to inter- or intramolecular interactions within the complex, the strength of the B-N bond is increased; and (4) conjugated imines.

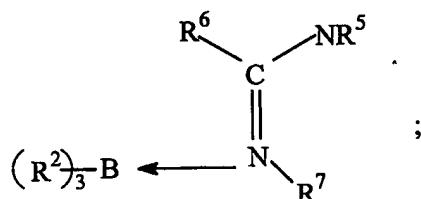
4. The fuel tank assembly of Claim 2 wherein
 500 the complex of the organoborane and the primary amine
 corresponds to the formula



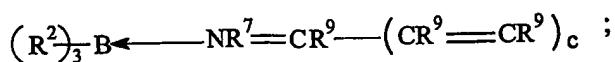
the organoborane heterocyclic amine complex corresponds
 to the formula



505 the organoborane amidine complex corresponds to the
 formula



and the organoborane conjugated imine complex
 corresponds to the formula



510

wherein:

B is boron; R¹ is separately in each occurrence
 hydrogen, a C₁₋₁₀ alkyl or C₃₋₁₀ cycloalkyl;

515 R² is separately in each occurrence a C₁₋₁₀ alkyl,
C₃₋₁₀ cycloalkyl or two or more of R² may combine to form a
cycloaliphatic ring structure;

520 R³ is separately in each occurrence hydrogen, a
C₁₋₁₀ alkyl or C₃₋₁₀ cycloalkyl; R⁴ is separately in each
occurrence hydrogen, C₁₋₁₀ alkyl, C₃₋₁₀ cycloalkyl, C₆₋₁₀ aryl
or alkaryl; R⁵, R⁶, and R⁷ are separately in each
occurrence hydrogen, C₁₋₁₀ alkyl, C₃₋₁₀ cycloalkyl, or two or
more of R⁵, R⁶ and R⁷ in any combination can combine to
form a ring structure which can be a single ring or a
multiple ring structure and the ring structure can include
525 one or more of nitrogen, oxygen or unsaturation in the
ring structure; R⁹ is independently in each occurrence
hydrogen, C₁₋₁₀ alkyl or C₃₋₁₀ cycloalkyl, Y, -(C(R⁹)₂-
(CR⁹=CR⁹)_c-Y or two or more of R⁹ can combine to form a ring
structure, or one or more of R⁹ can form a ring structure
530 with Y provided the ring structure is conjugated with
respect to the double bond of the imine nitrogen; X is a
hydrogen-bond accepting group with the proviso that where
the hydrogen bond accepting group is an amine it must be
secondary or tertiary;

535 Y is independently in each occurrence hydrogen,
N(R⁴)₂, OR⁴, C(O)OR⁴, a halogen or an alkylene group which
forms a cyclic ring with R⁷ or R⁹; Z is separately in each
occurrence oxygen or -NR⁴; a is separately in each
occurrence an integer of from 1 to 10; b is separately in
540 each occurrence 0 or 1, with the proviso that the sum of a
and b should be from 2 to 10; c is separately in each
occurrence an integer of from 1 to 10; x is separately in
each occurrence an integer of 1 to 10, with the proviso

that the total of all occurrences of x is from 2 to 10;
545 and y is separately in each occurrence 0 or 1.

5. The fuel tank assembly of Claim 2 wherein the organo borane/amine complex comprises an aliphatic heterocyclic amine which is a five or six-membered heterocyclic compound.

550 6. The fuel tank assembly of Claim 2 wherein the organo borane compound of the complex has three ligands selected from C₁₋₁₀ alkyl groups or phenyl groups, and the amine compound is selected from 1,6-diaminohexane, diethylamine, dibutylamine, diethylenetriamine, 555 dipropylenediamine, 1,3-propylenediamine, and 1,2-propylene-diamine.

560 7. The fuel tank assembly of Claim 2 wherein the organoborane compound of the complex has three ligands attached to the borane atom and which are selected from C₁₋₁₀ alkyl groups and phenyl and the amine compound is an alkanol amine or a diamine wherein the first amine group is a primary or secondary amine and the second amine is a primary amine.

565 8. The fuel tank assembly of Claim 2 wherein the amine compound of the complex is a polyoxyalkylene polyamine or a polyamine which is the reaction product of a diprimary amine and a compound having at least two groups which react with a primary amine.

570 9. The fuel tank assembly of Claim 1 wherein the low surface energy plastic material is a polyolefin.

10. The fuel tank assembly of Claim 9 wherein the polyolefin is selected from the group consisting of polyethylene, polypropylene and polytetrafluoroethylene.

11. The fuel tank assembly of Claim 1 wherein
575 the polymer having fuel barrier property is selected from
the group consisting of polyamides, fluoroelastomers,
polyacetal homopolymers and copolymers, sulfonated and
fluorinated HDPE, ethylene vinyl alcohol polymers and
copolymers, hydroxy-functionalized polyethers and
580 polyesters, and branched polyesters.

12. The fuel tank assembly of Claim 1 wherein
the fuel tank is a three-layer laminate structure
comprising two outer layers of a low energy surface
material and a core layer of a polymer having fuel barrier
585 property.

13. The fuel tank assembly of Claim 12 wherein
the low energy surface material is polyethylene and the
polymer having fuel barrier property is selected from the
group consisting of polyamides, fluoroelastomers,
590 polyacetal homopolymers and copolymers, sulfonated and
fluorinated HDPE, ethylene vinyl alcohol polymers and
copolymers, hydroxy-functionalized polyethers and
polyesters, and branched polyesters.

14. The fuel tank assembly of Claim 1 wherein
595 the tubular body has a first end and a second end, a
radially and outwardly extending fluted surface and a
raised surface spaced from the second end, and a bondline
extending from one side of the fluted surface to the
nearest side of the raised surface.

600 15. A fuel tank assembly comprising a fuel tank
having a wall with an outer surface and an inner surface,
a single or multi-walled fuel transport line having a
first open end and a second open end, the first open end
extending outwardly through an opening in the tank wall,

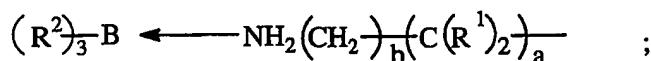
605 and the second open end extending inwardly into the tank
and bonded to the tank wall along the periphery of the
tank wall opening by an adhesive which bonds to low
surface energy plastic materials, the adhesive providing a
fuel vapor-tight seal at the interface between the fuel
610 transport line and tank wall opening, the fuel tank and
the fuel transport line comprising a polymer having fuel
barrier property.

16. The fuel tank assembly of Claim 15 wherein
the fuel transport line has a first end and a second end,
615 a radially and outwardly extending fluted surface and a
raised surface spaced from the second end, and a bondline
extending from one side of the fluted surface to the
nearest side of the raised surface.

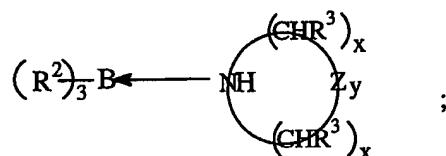
AMENDED CLAIMS

[received by the International Bureau on 17 October 2002 (17.10.02);
CLAIM 4 REPLACED]

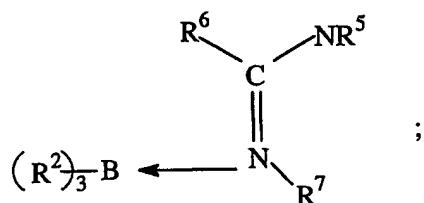
4. The fuel tank assembly of Claim 2 wherein
the complex of the organoborane and the primary amine
corresponds to the formula



the organoborane heterocyclic amine complex corresponds
to the formula

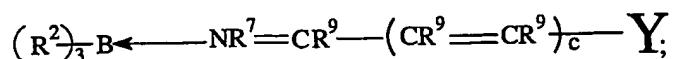


the organoborane amidine complex corresponds to the
formula



and the organoborane conjugated imine complex
corresponds to the formula

wherein:



B is boron; R¹ is separately in each occurrence
hydrogen, a C₁₋₁₀ alkyl or C₃₋₁₀ cycloalkyl;

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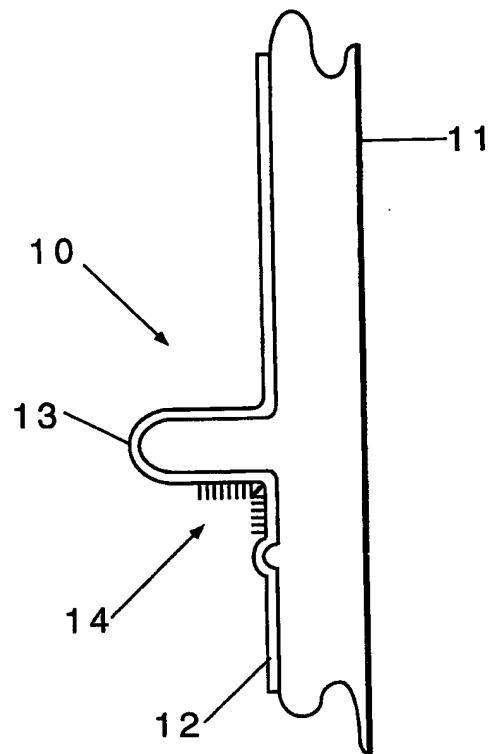


Fig. 1

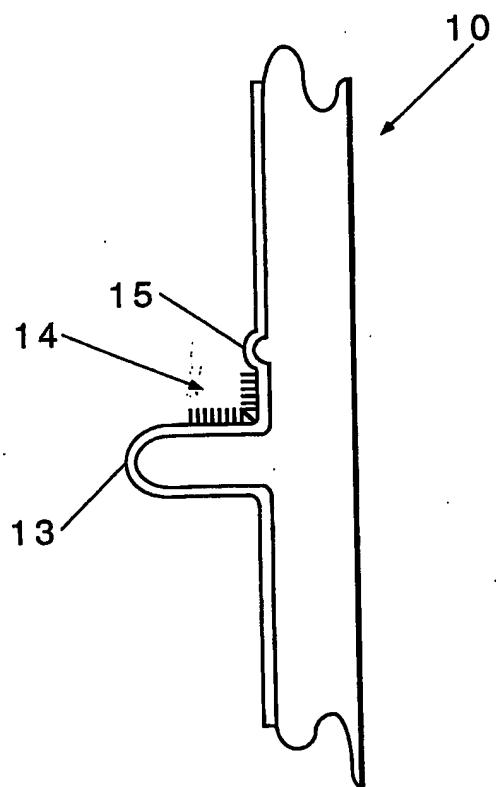


Fig. 2

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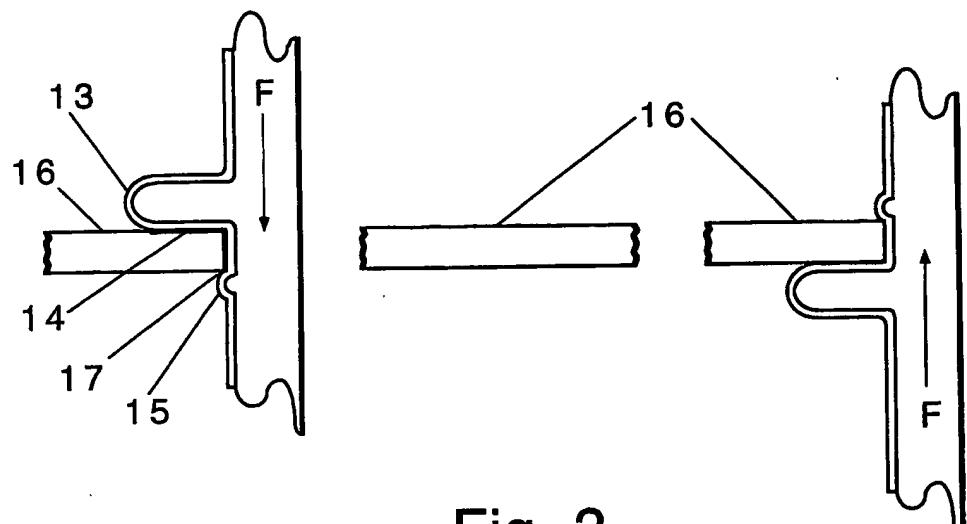


Fig. 3

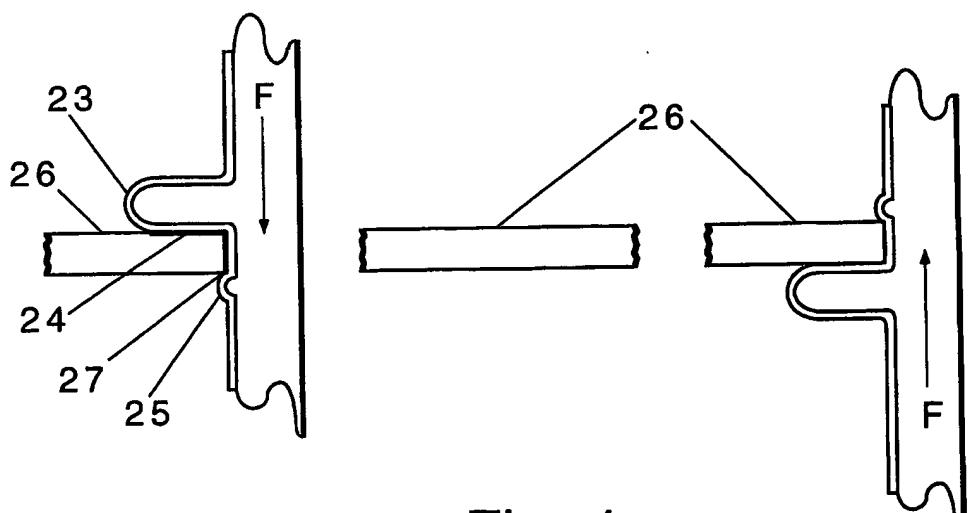


Fig. 4

INTERNATIONAL SEARCH REPORT

International Application No PCT/US 02/13975

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 B60K15/073 B60K15/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) IPC 7 B60K F16L C09J

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)
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PAJ, EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	PATENT ABSTRACTS OF JAPAN vol. 011, no. 296 (M-627), 25 September 1987 (1987-09-25) & JP 62 089584 A (TOYOTA MOTOR CORP; OTHERS: 02), 24 April 1987 (1987-04-24) abstract -----	1-16
A	DE 199 07 736 A (VOLKSWAGENWERK AG) 24 August 2000 (2000-08-24) claims 1-3; figures -----	1-16
A	US 5 690 780 A (KRASNOV DECEASED JURY N ET AL) 25 November 1997 (1997-11-25) cited in the application claim 1 -----	1-16

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Date of the actual completion of the International search

30 August 2002

Date of mailing of the International search report
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16/09/2002

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 02/13975

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
JP 62089584	A	24-04-1987	NONE		
DE 19907736	A	24-08-2000	DE	19907736 A1	24-08-2000
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